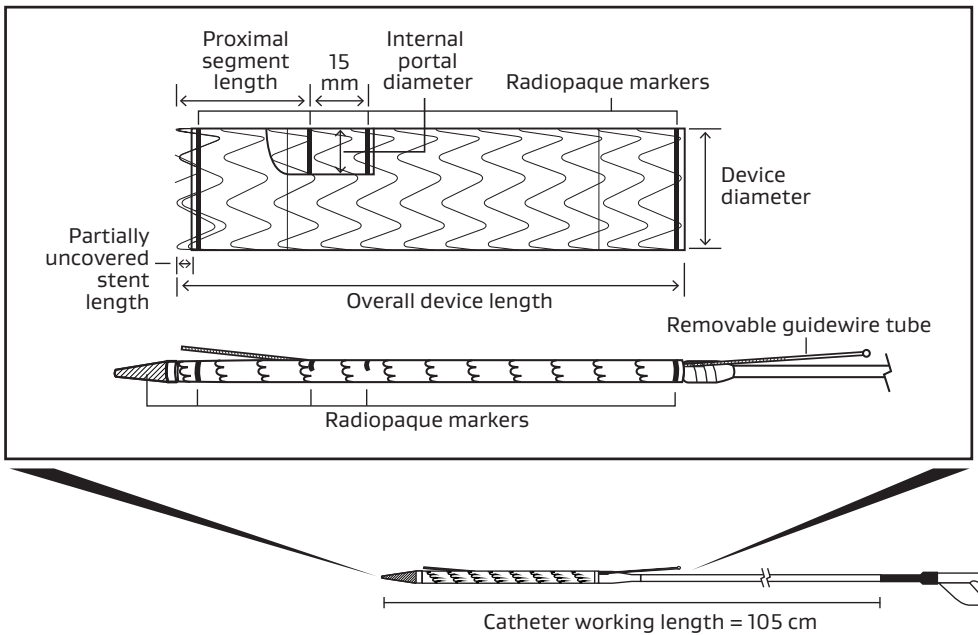
Together, improving life

GORE® TAG®
Thoracic Branch Endoprosthesis

DEVICE SIZING POCKET REFERENCE GUIDE

GORE® TAG® Thoracic Branch Endoprosthesis

Aortic Component and Aortic Component Delivery System



GORE® TAG® Thoracic Branch Endoprosthesis with 8 mm portal

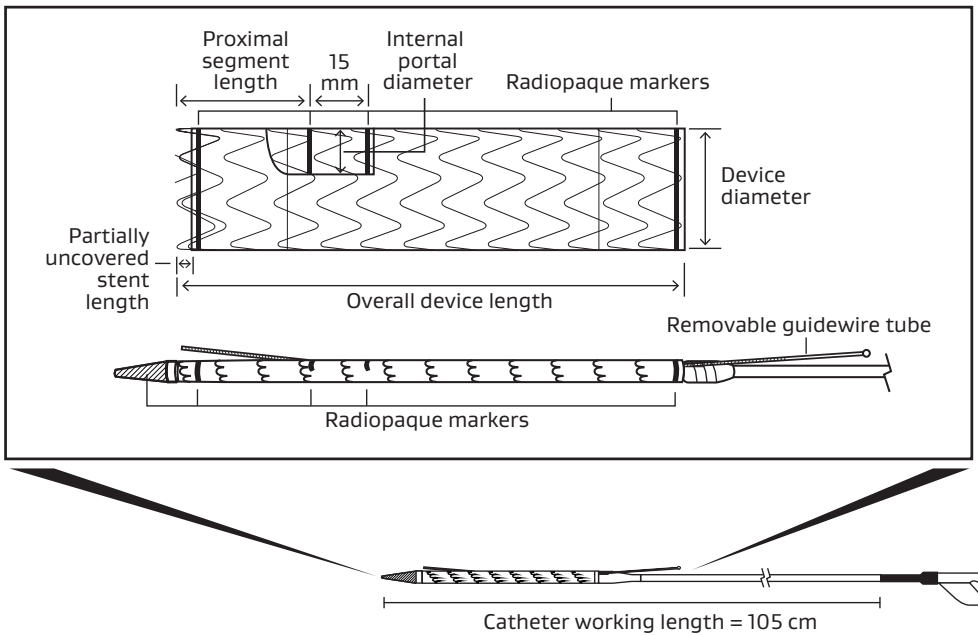
Aortic Component and Aortic Component Delivery System

Device dimensions and sizing requirements

Catalogue number	Device diameter (mm)	Intended aortic diameters (mm)	Internal portal diameter (mm)	Proximal segment length (mm)	Proximal covered length (mm)	Partially uncovered stent length (mm)	Overall device length (cm)	GORE® DRYSEAL Flex Introducer Sheath size (Fr)
TAC082110A	21	16–19.5	8	20	17	3	10	20
TAC082610A	26	19.5–24	8	20	16	4	10	20
TAC082615A	26	19.5–24	8	20	16	4	15	20
TAC082810A	28	22–26	8	20	16	4	10	22
TAC082815A	28	22–26	8	20	16	4	15	22
TAC082820A	28	22–26	8	20	16	4	20	22
TAC083115A	31	24–29	8	20	16	4	15	22
TAC083120A	31	24–29	8	20	16	4	20	22
TAC083415A	34	27–32	8	20	15	5	15	24
TAC083420A	34	27–32	8	20	15	5	20	24
TAC083715A	37	29–34	8	25	20	5	15	24
TAC083720A	37	29–34	8	25	20	5	20	24
TAC084015A	40	31–37	8	25	19	6	15	26
TAC084020A	40	31–37	8	25	19	6	20	26
TAC084515A	45	34–42	8	25	18.5	6.5	15	26
TAC084520A	45	34–42	8	25	18.5	6.5	20	26

GORE® TAG® Thoracic Branch Endoprosthesis

Aortic Component and Aortic Component Delivery System



GORE® TAG® Thoracic Branch Endoprosthesis with 12 mm portal

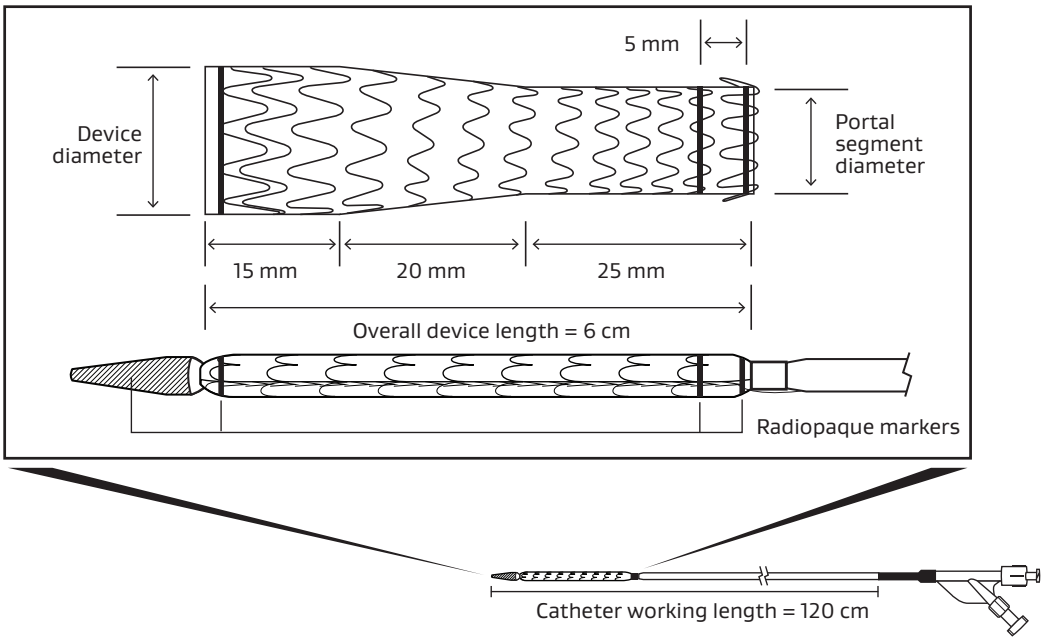
Aortic Component and Aortic Component Delivery System

Device dimensions and sizing requirements

Catalogue number	Device diameter (mm)	Intended aortic diameters (mm)	Internal portal diameter (mm)	Proximal segment length (mm)	Proximal covered length (mm)	Partially uncovered stent length (mm)	Overall device length (cm)	GORE® DRYSEAL Flex Introducer Sheath size (Fr)
TAC123115A	31	24–29	12	40	36	4	15	22
TAC123120A	31	24–29	12	40	36	4	20	22
TAC123415A	34	27–32	12	40	35	5	15	24
TAC123420A	34	27–32	12	40	35	5	20	24
TAC123715A	37	29–34	12	40	35	5	15	24
TAC123720A	37	29–34	12	40	35	5	20	24
TAC124015A	40	31–37	12	40	34	6	15	26
TAC124020A	40	31–37	12	40	34	6	20	26
TAC124515A	45	34–42	12	40	33.5	6.5	15	26
TAC124520A	45	34–42	12	40	33.5	6.5	20	26

GORE® TAG® Thoracic Branch Endoprosthesis

Side Branch Component and Side Branch Component Delivery System



GORE® TAG® Thoracic Branch Endoprosthesis

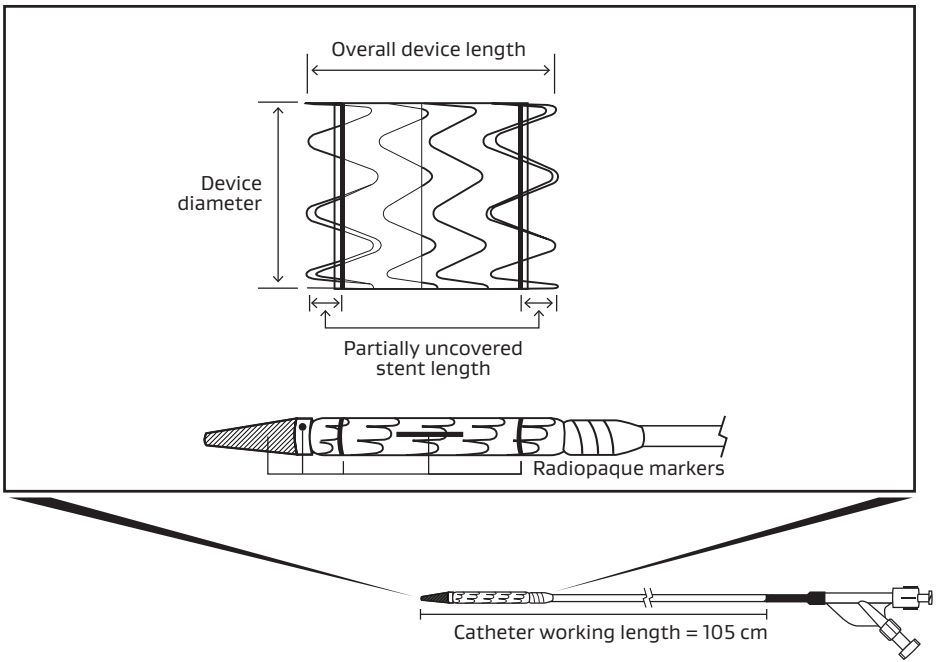
Side Branch Component and Side Branch Component Delivery System

Device dimensions and sizing requirements

Catalogue number	Device diameter (mm)	Intended branch vessel diameter (mm)	Portal segment diameter (mm)	Overall device length (cm)	Required branch vessel length (cm)	GORE® DRYSEAL Flex Introducer Sheath (Fr)
TSB080806A	8	6–7.5	8	6	3	14
TSB081006A	10	7.5–9	8	6	3	14
TSB081206A	12	9–11	8	6	3	14
TSB081506A	15	11–13	8	6	3	14
TSB081706A	17	13–15	8	6	3	14
TSB121506A	15	11–13	12	6	2.5	14
TSB121706A	17	13–15	12	6	2.5	14
TSB122006A	20	15–18	12	6	2.5	14

GORE® TAG® Thoracic Branch Endoprosthesis

Aortic Extender and Aortic Extender Delivery System



GORE® TAG® Thoracic Branch Endoprosthesis

Aortic Extender and Aortic Extender Delivery System

Device dimensions and sizing requirements

Catalogue number	Device diameter (mm)	Intended aortic diameters (mm)	Overall device length (cm)	Partially uncovered stent length (mm)	GORE® DRYSEAL Flex Introducer Sheath size (Fr)
TE2136A	21	16–19.5	3.6	3	20
TE2638A	26	19.5–24	3.8	4	20
TE2840A	28	22–26	4	4	22
TE3140A	31	24–29	4	4	22
TE3442A	34	27–32	4.2	5	24
TE3742A	37	29–34	4.2	5	24
TE4043A	40	31–37	4.3	6	26
TE4546A	45	34–42	4.6	6.5	26

GORE® TAG® Conformable Thoracic Stent Graft with ACTIVE CONTROL System

Straight devices

Catalogue number	Intended aortic diameter (mm)	Proximal diameter (mm)	Distal diameter (mm)	Endoprosthesis length (cm)	Recommended GORE® DRYSEAL Flex Introducer Sheath size (Fr)	Partially uncovered stent length (mm)
TGM212110	16–19.5	21	21	10	18	3
TGM262610	19.5–24	26	26	10	20	4
TGM282810	22–26	28	28	10	20	4
TGM282815	22–26	28	28	15	20	4
TGMR313110	24–29	31	31	10	20	4
TGMR313115	24–29	31	31	15	20	4
TGMR313120	24–29	31	31	20	20	4
TGM343410	27–32	34	34	10	22	5
TGM343415	27–32	34	34	15	22	5
TGM343420	27–32	34	34	20	22	5
TGMR373710	29–34	37	37	10	22	5
TGMR373715	29–34	37	37	15	22	5
TGMR373720	29–34	37	37	20	22	5
TGMR404010	31–37	40	40	10	22	6
TGMR404015	31–37	40	40	15	22	6
TGMR404020	31–37	40	40	20	22	6
TGM454510	34–42	45	45	10	24	6.5
TGM454515	34–42	45	45	15	24	6.5
TGM454520	34–42	45	45	20	24	6.5

GORE® TAG® Conformable Thoracic Stent Graft with ACTIVE CONTROL System

Tapered devices

Catalogue number[a]	Intended aortic diameter (mm)	Proximal diameter (mm)	Distal diameter (mm)	Endoprosthesis length (cm)	Recommended GORE® DRYSEAL Flex Introducer Sheath size (Fr)	Partially uncovered stent length (mm)
TGM262110	19.5–24/ 16–19.5	26	21	10	20	4
TGMR312610	24–29/ 19.5–24	31	26	10	20	4
TGM342815	27–32/ 22–26	34	28	15	22	5
TGMR373115	29–34/ 24–29	37	31	15	22	5
TGMR403415	31–37/ 27–32	40	34	15	22	6
TGM453715	34–42/ 29–34	45	37	15	24	6.5

a. Some sizes not available in all markets.

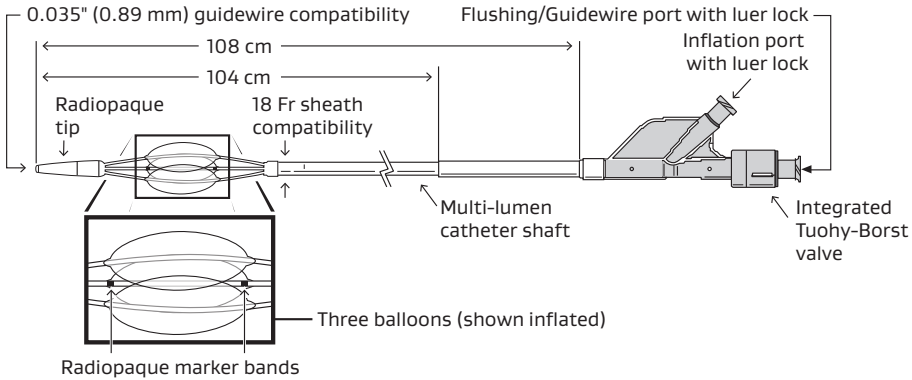
GORE® DRYSEAL Flex Introducer Sheath

GORE® DRYSEAL Flex Introducer Sheath

Catalogue number[b]	Sheath size (Fr)	Outer diameter (mm)	Sheath working length (cm)
GDSF1833	18	6.7	33
GDSF1865	18	6.7	65
GDSF2033	20	7.5	33
GDSF2065	20	7.5	65
GDSF2233	22	8.2	33
GDSF2265	22	8.2	65
GDSF2433	24	8.8	33
GDSF2465	24	8.8	65
GDSF2633	26	9.5	33
GDSF2665	26	9.5	65

b. GDSF catalogue numbers are not available in all regions; use DSF catalogue numbers instead where appropriate.

GORE® Tri-Lobe Balloon Catheter



Device dimensions and sizing requirements

U.S. catalogue number[c]	Inner vessel diameter size	Working length (cm)	Size (Fr)
BCM1634	Aortic diameters 16–32 mm	104	18
BCL2645	Aortic diameters 26–42 mm	104	18
TBCM1634	Aortic diameters 16–32 mm	104	18
TBCL2648	Aortic diameters 26–48 mm	104	18

Canada catalogue number	Inner vessel diameter size	Working length (cm)	Size (Fr)
BCM1634	Aortic diameters 16–32 mm	104	18
BCL2645	Aortic diameters 26–48 mm	104	18

[c] Products listed may not be available in all markets. For accounts where TBCM1634 and TBCL2648 are not available, BCM1634 and BCL2645 may be substituted.

Consult Instructions for Use
eifu.goremedical.com

Refer to *Instructions for Use* at eifu.goremedical.com for a complete description of all applicable indications, warnings, precautions and contraindications for the markets where this product is available. $\text{R}_{\text{X Only}}$

GORE® DRYSEAL Flex Introducer Sheath

INDICATIONS FOR USE IN THE U.S.: The GORE® DRYSEAL Flex Introducer Sheath is intended to be inserted in the vasculature to provide a conduit for the insertion of endovascular devices while minimizing blood loss associated with such insertions. **CONTRAINDICATIONS:** There are no known contraindications for this device.

GORE® TAG® Conformable Thoracic Stent Graft

INDICATIONS FOR USE IN THE U.S.: The GORE® TAG® Conformable Thoracic Stent Graft is intended for endovascular repair of all lesions of the descending thoracic aorta, including: isolated lesions in patients who have appropriate anatomy, including: adequate iliac/femoral access, aortic inner diameter in the range of 16-42 mm, ≥ 20 mm non-aneurysmal aorta proximal and distal to the lesion; Type B dissections in patients who have appropriate anatomy, including: adequate iliac/femoral access, ≥ 20 mm landing zone proximal to the primary entry tear; proximal extent of the landing zone must not be dissected, diameter at proximal extent of proximal landing zone in the range of 16-42 mm. **CONTRAINDICATIONS:** Patients with known sensitivities or allergies to the device materials; patients who have a condition that threatens to infect the graft.

GORE® TAG® Thoracic Branch Endoprosthesis

INDICATIONS FOR USE: The GORE® TAG® Thoracic Branch Endoprosthesis is indicated for endovascular repair of lesions of the aortic arch and descending thoracic aorta, while maintaining flow into a single aortic arch branch vessel in patients who have: **Adequate iliac/femoral access; Proximal Aortic Landing Zones:** For Isolated Lesion Patients: Proximal landing zone cannot be aneurysmal, dissected, heavily calcified or heavily thrombosed; For Dissection Patients: Primary entry tear must be distal to the target branch vessel and the proximal extent of the landing zone must not be dissected; Aortic inner diameter range 16–42 mm; Proximal segment length (length from distal edge of target branch vessel to the midpoint of any proximal branch vessel) of at least 2.0–4.0 cm, depending on Aortic

Component selection; Proximal covered length (measured from distal edge of target branch vessel to the distal edge of any proximal branch vessel) of at least 15–36 mm, depending on Aortic Component selection; For patients with prior ascending aorta or aortic arch repair with surgical graft: at least 2 cm landing zone proximal to the distal anastomosis; **Target Branch Vessel Landing Zone:** Landing zone cannot be aneurysmal, dissected, heavily thrombosed and severely tortuous (180 degree turn within the treated length); Target branch vessel inner diameter of 6–18 mm, depending on Side Branch Portal diameter selected; Target branch vessel minimum length of 2.5–3.0 cm, depending on Side Branch Portal diameter selected. **Distal Landing Zone (Isolated Lesion Patients only):** Outer curve length must be ≥ 2 cm proximal to celiac artery; Aortic inner diameter range 16–42 mm; Cannot be aneurysmal, dissected, heavily calcified or heavily thrombosed; Native Aorta or previously placed GORE® TAG® Conformable Thoracic Stent Graft. **CONTRAINDICATIONS:** The GORE® TAG® Thoracic Branch Endoprosthesis is contraindicated in: Patients with known sensitivities or allergies to the device materials [ePTFE (polytetrafluoroethylene), FEP (fluoroethylpropylene), Nitinol (nickel, titanium), Gold, SB Component only - Heparin (CBAS® Heparin Surface)]; Patients who have a condition that threatens to infect the graft; Patients with known hypersensitivity to heparin, including those patients who have had a previous incident of Heparin-Induced Thrombocytopenia (HIT) type II.

GORE® Tri-Lobe Balloon Catheter

INDICATIONS FOR USE IN CANADA: The GORE® Tri-Lobe Balloon Catheter is intended to facilitate in the endovascular repair of the thoracic or abdominal aorta due to lesions including aneurysms, dissections, trauma, and penetrating aortic ulcers. **CONTRAINDICATIONS:** There are no known contraindications."

INDICATIONS FOR USE IN THE U.S.: The GORE® Tri-Lobe Balloon Catheter is intended to assist in the dilatation of self-expanding endoprostheses in large diameter vessels. **CONTRAINDICATIONS:** There are no known contraindications.

Products listed may not be available in all markets.

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